

LYME IBGM, SAMPLE1

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DOB:

Gender: Accession:

Patient ID:

Physician: IGENEX, INC. Client: 1

Date of Collection: 11/14/2017 07:00

Sample Received 11/14/2017 17:20

Report Printed: 11/14/2017 17:26

TEST	SPECIMEN	RESULT	REFERENCE RANGE	UNITS
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Lyme ImmunoBlot IgG (B. burgdorferi sensu lato)

IGENEX CRITERIA:

Positive= Presence of 2 or more of the following bands:
23, 31, 34, 39, 41 and 93 kDa

Negative= Any profile that does not meet positive criteria

CDC/NYS CRITERIA:

Positive= Presence of 5 or more of the following bands:
18, 23, 28, 30, 39, 41, 45, 58, 66, 93 kDa

Negative= Any profile that does not meet positive criteria

Band Intensity: (+ to +++)positive, (Ind)indeterminate, (-)negative

18 kDa	Serum	+
23 kDa	Serum	-
28 kDa	Serum	-
30 kDa	Serum	-
31 kDa	Serum	+
34 kDa	Serum	+
39 kDa	Serum	+
41 kDa	Serum	+
45 kDa	Serum	+
58 kDa	Serum	+
66 kDa	Serum	+
93 kDa	Serum	+
IGeneX Criteria Result	Serum	Pos
CDC/NYS Criteria Result	Serum	Pos

* Lyme ImmunoBlot IgG detects antibodies to *B. burgdorferi* B31, *B. burgdorferi* 297, *B. mayonii*, *B. californiensis*, *B. spielmanii*, *B. valaisiana*, *B. atzeli* and *B. garinii*

Approval	Initial	Date
Lab Director	<i>JS</i>	11-14-17
Lab Manager	<i>AL</i>	11-14-17
CLS	<i>SP</i>	11-14-17

End of Report

*Testing performed at IGeneX 795 San Antonio Road Palo Alto CA 94303 (800) 832-3200

Diagnosis should not be based on laboratory results alone. Results should be interpreted in conjunction with clinical symptoms and patient history.

NOTE: Western Blots, ImmunoBlots, Lyme Dot Blot, Epitope, PCR, IFA, FISH, C. pneumoniae IgG/IgA, CD57, IGXSpot, Lyme Array Test - These tests were developed and their performance characteristics determined by IGeneX, Inc. They have not been cleared or approved by the FDA. The FDA has determined that such approval is not necessary. These tests are used for clinical purposes and should not be regarded as investigational or for research. IGeneX, Inc. is licensed by CMS and NYS to perform high complexity clinical laboratory testing.